AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Currently Amended) A method of preventing or treating carnitine deficiency in chronic uremic patients undergoing periodic dialysis comprising administering to the patient at the conclusion of the dialysis an effective amount of into a venous return line after a dialysis session from about 10 to about 20 mg/kg body weight of carnitine, calculated as L-carnitine or a pharmaceutically acceptable salt thereof wherein the treatment is repeated twice a week every 44 hours, then after 68 hours.
 - 2.-5. Canceled.
- 6. (Currently Amended) The method of claim 51, wherein the treatment is continued for3-4 weeks, monitoring pre-dialytic levels of carnitine.
- 7. (Original) The method of claim 6, wherein pre-dialytic levels of carnitine are monitored.
- 8. (Original) The method of claim 7, wherein pre-dialytic levels of carnitine are equal to lower than $40\text{-}50\mu M$.
- 9. (Currently Amended) The method of claim 41, wherein a maintenance dosage is provided, administering a dose of 5 mg/kg of carnitine.
 - 10. Canceled.
- 11. (Original) The method of claim 1, wherein carnitine fumarate is the pharmaceutically acceptable salt.
- 12. (Original) The method of claim 11, wherein the patient is affected by hypervolemic heart.

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- 13. (Original) The method of claim 11, wherein the patient is affected by diabetes.
- 14. (Currently Amended) A method of preventing or treating carnitine deficiency in chronic uremic patients undergoing periodic dialysis comprising administering to the patient at the conclusion of the dialysis into a venous return line after a dialysis session an amount of L-carnitine or of a pharmaceutically acceptable salt thereof effective to restore a level of carnitine in the patentpatient to a pre-dialytic level, and thereafter reducing the amount of carnitine administered to a level sufficient to maintain carnitine levels to the pre-dialytic level.
- 15. (Currently Amended) A method of preventing or treating carnitine deficiency in chronic uremic patients undergoing periodic dialysis comprising administering to the patient at the conclusion of the dialysis into a venous return line after a dialysis session from about 10 to about 20 mg/kg body weight of carnitine, calculated as L-carnitine, or of a pharmaceutically acceptable salt thereof to restore a level of carnitine in the patent to a pre-dialytic level, and thereafter reducing the amount of carnitine administered to a level sufficient to maintain carnitine levels to the pre-dialytic level.
- 16. (Original) The method of claim 15 wherein the treatment to achieve pre-dialytic levels is on a weekly basis repeated twice a week every 44 hours, then after 68 hours.
 - 17. (Original) The method of claim 16, wherein the treatment is continued for 3-4 weeks.
- 18. (Original) The method of claim 14 or 15, wherein pre-dialytic levels of carnitine are equal or lower than $40\text{-}50\mu\text{M}$.
- 19. (Currently Amended) The method of claim 14 of 15, wherein a maintenance dosage of about 5 mg/kg of carnitine is administered.